DEXTROMETHORPHAN POLISTIREX EXTENDED RELEASE- dextromethorphan polistirex suspension Bryant Ranch Prepack

Perrigo Dextromethorphan Polistirex Extended-Release Oral Suspension Drug Facts

Active ingredient (in each 5 mL)

Dextromethorphan polistirex equivalent to 30 mg dextromethorphan hydrobromide

Purpose

Cough suppressant

Uses

temporarily relieves

- cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
- the impulse to cough to help you get to sleep

Warnings

Do not use

if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Allergy Alert: Contains sodium metabisulfite, a sulfite that may cause allergic-type reactions.

Ask a doctor before use if you have

- chronic cough that lasts as occurs with smoking, asthma or emphysema
- cough that occurs with too much phlegm (mucus)

Stop use and ask a doctor if

- side effects occur. You may report side effects to FDA at 1-800-FDA-1088.
- cough lasts more than 7 days, cough comes back, or occurs with fever, rash or headache that lasts. These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- shake bottle well before use
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by doctor

| adults and children 12 years of age and over | 10 mL every 12 hours, not to exceed 20 mL in 24 hours |
|--|---|
| children 6 to under 12 years of age | 5 mL every 12 hours, not to exceed 10 mL in 24 hours |
| children 4 to under 6 years of age | 2.5 mL every 12 hours, not to exceed 5 mL in 24 hours |
| children under 4 years of age | do not use |

Other information

- each 5 mL contains: sodium 5 mg
- store at 20° to 25°C (68° to 77°F)
- dosing cup provided

Inactive ingredients

D&C Red #30 aluminum lake, D&C Yellow #10 aluminum lake, flavor, glycerin, high fructose corn syrup, methylparaben, polysorbate 80, polyvinyl acetate, povidone, propylparaben, purified water, sodium metabisulfite, sodium polystyrene sulfonate, sucrose, tartaric acid, tragacanth gum, triacetin, xanthan gum

Questions or comments?

1-800-719-9260

HOW SUPPLIED

Dextromethorphan Polistirex Extended-Release Oral Suspension

NDC 72162-2281-2: 89 mL in a BOTTLE

Repackaged/Relabeled by:

Bryant Ranch Prepack, Inc.

Dextromethorphan Polistirex Extended-Release Oral Suspension



SECT

Drug Facts
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Uses
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Do not use if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drug for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product. Allerny Alert: Contains sodium metablisuffice, a suffer that may cause altergic-type reactions. Ask a doctor before use if you have -chronic cough that lasts as occus with smoking, ashtma or emphysema-cough that occurs with or or much philegrin (mucus) Slov use and ask a doctor if side effects occur. You may report side effects of extending mucus) Slov use and ask a doctor if side effects occur. You may report side effects of ever, rash or headache that lasts. These could be signs of a serious condition. If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poisso Control Center right way (1-800-222-1222).

Other Information

each 5 mL contains: sodium 5 mg *store at 20° to 25°C (68° to 77°F) *dosing cup provided. *1
 Hour Cough Relief *Day or Night *Alcohol-free *Orange Flavored Liquid.

Directions
-shake bottle

•shake bottle well before use - measure only with dosing cup provided -do not use dosing cup with other products - dose as follows or as directed by doctor, (adults and other 12 years of age and over, 10 mt. every 12 hours, not to exceed 20 mt. in 24 hours).(children 6 to under 12 years of age. 2.5 mt. every 12 hours, not to exceed 70 mt. in 24 hours). (children 4 to under 6 years of age, 2.5 mt. every 12 hours, not to exceed 5 mt. in 24 hours). (children 4 years of age, 30 not use). Inactive Ingredients

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NDC 72162-2281-2

Dextromethorphan Polistirex Extended-Release Oral Suspension

30 mg/5 mL

BRP

3 FL OZ (89 mL)

Relabeled by:

Bryant Ranch Prepack, Inc. Burbank, CA 91504 USA



Extended Label

Package Insert

| SHAKE WELL Measure only with | SING . BEFORE USE. dosing cup provided. up with other products. |
|---------------------------------|--|
| Age (yr) | Dose |
| 12 years to adult | 10 mL EVERY 12 HOURS |
| 6 to under 12 | 5 mL EVERY 12 HOURS |
| 4 to under 6 | 2.5 mL EVERY 12 HOURS |
| Under 4 | Do not use |

DEXTROMETHORPHAN POLISTIREX EXTENDED RELEASE

dextromethorphan polistirex suspension

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72162-2281(NDC:45802-433)

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)
(DEXTROMETHORPHAN - UNII:7355X3ROTS)

DEXTROMETHORPHAN | 30 mg in 5 mL

| Inactive Ingredients | | |
|--|--|--|
| Ingredient Name Streng | | |
| POLISTIREX (UNII: 5H9W9GTW27) | | |
| D&C RED NO. 30 (UNII: 2S42T2808B) | | |
| D&C YELLOW NO. 10 (UNII: 35SW5USQ3G) | | |

| Product Characteristics | | |
|-------------------------|--------|--------------|
| Color | ORANGE | Score |
| Shape | | Size |
| Flavor | ORANGE | Imprint Code |
| Contains | | |

| Packaging | | | |
|------------------------|--|-------------------------|-----------------------|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 NDC:72162- 2281-2 | 1 in 1 CARTON | 04/30/2024 | |
| 1 | 89 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |

| Marketing Information | | | |
|---|------------|-------------------------|-----------------------|
| Marketing Application Number or Monograph Category Citation | | Marketing Start Date | Marketing End Date |
| ANDA | ANDA091135 | 09/10/2012 | |
| | | | |

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

| Establishment | | | |
|----------------------|---------|-----------|---|
| Name | Address | ID/FEI | Business Operations |
| Bryant Ranch Prepack | | 171714327 | REPACK(72162-2281), RELABEL(72162-2281) |

Revised: 4/2025 Bryant Ranch Prepack